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L17: Entry 2 of 28

File: USPT

Mar 12, 2002

US-PAT-NO: 6355229

DOCUMENT-IDENTIFIER: US 6355229 B1

TITLE: Oral composition containing cetylpyridinium chloride and guar hydroxypropyltrimonium chloride and method of using the same

DATE-ISSUED: March 12, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Adamy; Steven T.	Hamilton	NJ		

US-CL-CURRENT: 424/54; 424/435, 424/440, 424/464, 424/48, 424/49

CLAIMS:

What is claimed is:

1. An oral composition comprising:

a) an antibacterial effective amount of cetylpyridinium chloride;b) an effective amount of guar hydroxypropyltrimonium chloride, sufficient to bind to compounds which undesirably bind to cetylpyridinium chloride thereby enabling the cetylpyridinium chloride to effectively bind to tooth surfaces and perform an antibacterial function; and

c) an orally acceptable carrier.

2. The oral composition of claim 1, wherein the orally acceptable carrier is selected from the group consisting of water, saline, alcohol, glycerin, oil and mixtures thereof.

3. The oral composition of claim 1, wherein the oral composition is in a form selected from the group consisting of a mouthwash, a dentifrice, a chewing gum, and a lozenge.4. The oral composition of claim 1, the amount of cetylpyridinium chloride is present from about 0.01 to 1.0% by weight based on the total weight of the oral composition.

5. The oral composition of claim 1, wherein the amount of guar hydroxypropyltrimonium chloride is present from about 0.1 to 3.0% by weight based on the total weight of the oral composition.

6. The oral composition of claim 3 wherein the dentifrice is a toothpaste.

7. The oral composition of claim 6 further comprising at least one material selected from the group consisting of thickening agents, whiteners, flavorants, humectants, desensitizing agents, abrasive agents, alkali metal bicarbonate salts, and fluoride supplying compounds.

8. The oral composition of claim 7 wherein the abrasive agents are selected from the group consisting of sodium metaphosphate, potassium metaphosphate,

tricalcium phosphate, dicalcium phosphate dihydrate, anhydrous dicalcium phosphate, calcium pyrophosphate, zinc orthophosphate, alumina, hydrated alumina, aluminum silicate, bentonite, calcium carbonate, and sodium bicarbonate.

9. The oral composition of claim 1 further comprising at least one sweetening agent.

10. The oral composition of claim 9 comprising at least one high potency sweetening agent.

11. The oral composition of claim 1 further comprising at least one additional antibacterial agent.

12. The oral composition of claim 11 wherein the at least one additional antibacterial agent is present in an amount of from about 0.1 to 2% by weight based on the total weight of the oral composition.

13. The oral composition of claim 7 wherein the abrasive agent is present in an amount of from about 0.5 to 70% by weight.

14. The oral composition of claim 7 wherein the fluoride supplying compound is present in an amount sufficient to deliver from about 100 to 5,000 ppm of available fluoride based on the composition.

15. The oral composition of claim 7 wherein the alkali metal bicarbonate salts are present in an amount of up to about 75% by weight based on the total weight of the oral composition.

16. The oral composition of claim 15 wherein the amount of the alkali metal bicarbonate salts are present in an amount of from about 5 to 40% by weight.

17. The oral composition of claim 7 wherein the thickening agent is present in an amount of from about 0.1 to 3.0% by weight based on the total weight of the composition.

18. The oral composition of claim 1 wherein the orally acceptable carrier is present in an amount of from about 20 to 99% by weight based on the total weight of the composition.

19. The oral composition of claim 7 wherein the humectant is present in an amount of from about 1 to 50% by weight based on the total weight of the composition.

20. A method of reducing the presence of microorganisms in an oral cavity of a warm-blooded animal, said method comprising administering to the oral cavity an effective amount of the oral composition of claim 1.

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NAME	CITY	STATE	ZIP CODE	COUNTRY
Adamy; Steven T.	Hamilton	NJ		

ASSIGNEE-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY	TYPE CODE
Church & Dwight Co., Inc.	Princeton	NJ			02

APPL-NO: 09/ 893766 [PALM]

DATE FILED: June 27, 2001

INT-CL: [07] A61 K 7/16, A61 K 7/22, A61 K 9/20, A61 K 9/68

US-CL-ISSUED: 424/54; 424/48, 424/49, 424/435, 424/440, 424/464

US-CL-CURRENT: 424/54; 424/435, 424/440, 424/464, 424/48, 424/49

FIELD-OF-SEARCH: 424/48, 424/49, 424/54, 424/435, 424/440, 424/464

PRIOR-ART-DISCLOSED:

U.S. PATENT DOCUMENTS

Search Selected

Search ALL

PAT-NO	ISSUE-DATE	PATENTEE-NAME	US-CL
<input type="checkbox"/> 4389394	June 1983	Drucker	
<input type="checkbox"/> 4405654	September 1983	Lee	
<input type="checkbox"/> 4585650	April 1986	Newberry et al.	424/73
<input type="checkbox"/> 4839158	June 1989	Michaels	
<input type="checkbox"/> 4847076	July 1989	Deshpande et al.	424/70.13
<input type="checkbox"/> 4971797	November 1990	Cherukuri	
<input type="checkbox"/> 5013763	May 1991	Tubesing et al.	424/195.18
<input type="checkbox"/> 5037643	August 1991	Green	424/78.23
<input type="checkbox"/> 5076953	December 1991	Jordan et al.	510/151
<input type="checkbox"/> 5085857	February 1992	Reid et al.	424/70.12
<input type="checkbox"/> 5152914	October 1992	Forster et al.	510/122
<input type="checkbox"/> 5158763	October 1992	Gaffar et al.	
<input type="checkbox"/> 5370881	December 1994	Fuisz	
<input type="checkbox"/> 5374638	December 1994	Hauschild	
<input type="checkbox"/> 5376360	December 1994	Domke et al.	
<input type="checkbox"/> 5523079	June 1996	Gough	424/70.11
<input type="checkbox"/> 5622689	April 1997	Lukacovic	
<input type="checkbox"/> 5801116	September 1998	Cottrell et al.	502/404
<input type="checkbox"/> 5849268	December 1998	Lukacovic	
<input type="checkbox"/> 5849271	December 1998	Lukacovic	
<input type="checkbox"/> 5948390	September 1999	Nelson et al.	
<input type="checkbox"/> 6274128	August 2001	Bergmann et al.	424/70.1

FOREIGN PATENT DOCUMENTS

FOREIGN-PAT-NO	PUBN-DATE	COUNTRY	US-CL
0 009 408	April 1980	EP	
0 422 803	April 1991	EP	
0 480 811	April 1992	EP	
0 507 598	October 1992	EP	
0 920 857	June 1999	EP	
WO 90/15592	June 1990	WO	
WO 97/46217	December 1997	WO	

ART-UNIT: 1614

PRIMARY-EXAMINER: Rose; Shep K.

ATTY-AGENT-FIRM: Fishman; Irving M.

ABSTRACT:

An oral composition comprises an antibacterial effective amount of cetylpyridinium chloride, an effective amount of guar hydroxypropyltrimonium chloride, sufficient to bind to compounds which undesirably bind to cetylpyridinium chloride thereby enabling the cetylpyridinium chloride to effectively bind to tooth surfaces and perform an antibacterial function, and an orally acceptable carrier.

20 Claims, 0 Drawing figures

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US-CL-CURRENT: 424/54; 424/435, 424/440, 424/464, 424/48, 424/49

FIELD-OF-SEARCH: 424/48, 424/49, 424/54, 424/435, 424/440, 424/464

PRIOR-ART-DISCLOSED:

U.S. PATENT DOCUMENTS

Search Selected

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cetylpyridinium chloride
+
zinc chloride

	PAT-NO	ISSUE-DATE	PATENTEE-NAME	US-CL
<input type="checkbox"/>	<u>4389394</u>	June 1983	Drucker	
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<input type="checkbox"/>	<u>5374638</u>	December 1994	Hauschild	
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<input type="checkbox"/>	<u>5523079</u>	June 1996	Gough	424/70.11
<input type="checkbox"/>	<u>5622689</u>	April 1997	Lukacovic	
<input type="checkbox"/>	<u>5801116</u>	September 1998	Cottrell et al.	502/404
<input type="checkbox"/>	<u>5849268</u>	December 1998	Lukacovic	
<input type="checkbox"/>	<u>5849271</u>	December 1998	Lukacovic	
<input type="checkbox"/>	<u>5948390</u>	September 1999	Nelson et al.	
<input type="checkbox"/>	<u>6274128</u>	August 2001	Bergmann et al.	424/70.1

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An oral composition comprises an antibacterial effective amount of cetylpyridinium chloride, an effective amount of guar hydroxypropyltrimonium chloride, sufficient to bind to compounds which undesirably bind to cetylpyridinium chloride thereby enabling the cetylpyridinium chloride to effectively bind to tooth surfaces and perform an antibacterial function, and an orally acceptable carrier.

20 Claims, 0 Drawing figures

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TITLE: Oral composition containing cetylpyridinium chloride and guar hydroxypropyltrimonium chloride and method of using the sameAbstract Text (1):

An oral composition comprises an antibacterial effective amount of cetylpyridinium chloride, an effective amount of guar hydroxypropyltrimonium chloride, sufficient to bind to compounds which undesirably bind to cetylpyridinium chloride thereby enabling the cetylpyridinium chloride to effectively bind to tooth surfaces and perform an antibacterial function, and an orally acceptable carrier.

Brief Summary Text (2):

The present invention relates to oral compositions containing cetylpyridinium chloride as an active antibacterial agent which may be used to inhibit formation of plaque, oral malodor, gingivitis, periodontal disease and the like. The oral composition contains the active agent and an effective amount of guar hydroxypropyltrimonium chloride which enables the active agent to more effectively bind to tooth surfaces to perform its antibacterial function.

Brief Summary Text (4):

Cetylpyridinium chloride (CPC) is well known as an antibacterial agent especially for the inhibition of plaque formation. This antibacterial agent has been used in commercial mouthwash products such as Scope.RTM. and Cepacol.RTM. and several other types of oral care products. The environment of such commercial mouthwash products enables cetylpyridinium chloride to freely contact those oral surfaces which may harbor unwanted microorganisms. These microorganisms contribute to both the initiation and progression of gingivitis, plaque, periodontal disease, and/or breath malodor in the oral cavity of warm-blooded animals. Such conditions are usually treated by reducing the presence of the microorganisms in the oral cavity through the use of dental care products containing antibacterial agents including cetylpyridinium chloride.

Brief Summary Text (5):

The antibacterial activity of cetylpyridinium chloride is, without being bound to the theory, believed to be linked to the cationic charge of its amine group. Thus, cetylpyridinium chloride is attracted to and binds to negatively-charged protein moieties on the cell membrane or cell wall of the microorganism and to tooth surfaces which are also typically negatively charged. The resulting attachment to microorganisms disrupts the cell wall structure causing leakage of the intracellular fluids, eventually killing the associated microorganism. However, cetylpyridinium chloride is generally not effective in many systems because of its tendency to complex with components that carry a negative charge. When bound to negatively charged particles in this manner, cetylpyridinium chloride is unavailable for effective contact with tooth surfaces and microorganisms, thereby rendering the active agent ineffective for its intended purpose.

Brief Summary Text (6):

For this reason, cetylpyridinium chloride has not been totally effective in typical oral care products for the treatment and/or prevention of gingivitis, plaque, periodontal disease, and/or breath malodor. For example, toothpaste compositions typically include anionic surfactants and artificial sweetening agents. These components of toothpaste compositions typically bind to cetylpyridinium chloride and thereby render the same ineffective or substantially less effective as an

Brief Summary Text (7):

Brief Summary Text. (9):

Brief Summary Text (10):

Brief Summary Text (11):

Brief Summary Text (15):

Brief Summary Text (16):

Brief Summary Text (17):

Brief Summary Text (18):

The oral composition of the present invention may be in the form of a solution such as a mouthwash, a denture cleanser, a gargle or the like, or may be in the form of a

semi-solid or solid such as a toothpaste, a gel dentifrice, a dental powder, a denture cleansing tablet, a chewing gum, or a solid lozenge or the like.

Brief Summary Text (19):

Cetylpyridinium chloride is cationic and therefore is attracted to negative surfaces and moieties. Tooth surfaces typically have a negative charge and therefore there is a natural attraction of cetylpyridinium chloride for tooth surfaces. However, many conventional oral care products contain components, including those found in toothpaste, which are anionic. Such negatively charged components bind to cetylpyridinium chloride and therefore make the antibacterial agent less available for binding to tooth surfaces and microorganisms.

Brief Summary Text (20):

In accordance with one aspect of the present invention, it has been discovered that the employment of guar hydroxypropyltrimonium chloride at least reduces and may substantially eliminate the ability of such compounds to bind to and diminish the antibacterial activity of cetylpyridinium chloride.

Brief Summary Text (21):

By way of example, certain sweetening agents known for use in oral compositions, such as saccharin, acesulfame, and cyclamate have been found to bind to and thereby inhibit the antibacterial activity of cetylpyridinium chloride when compared to compositions which do not contain such particular sweetening agents. In the oral composition of the present invention, the presence of typical sweetening agents does not adversely affect the antibacterial activity of cetylpyridinium chloride when guar hydroxypropyltrimonium chloride is present in effective amounts.

Brief Summary Text (23):

The present invention also provides a method of reducing the presence of microorganisms in a oral cavity comprising administering to the oral cavity an effective amount of an oral composition comprising an antibacterial effective amount of cetylpyridinium chloride and an effective amount of guar hydroxypropyltrimonium chloride. The present oral composition enables cetylpyridinium chloride to effectively bind to the tooth surfaces to perform an antibacterial function.

Brief Summary Text (24):

Guar hydroxypropyltrimonium chloride is incorporated into the oral composition of the present invention in an effective amount which is sufficient to bind to compounds which have the ability to bind to cetylpyridinium chloride. If a sufficient amount of such compounds can no longer bind to the antibacterial agent, then a sufficient amount of cetylpyridinium chloride will be available to perform its antibacterial function. Generally, guar hydroxypropyltrimonium chloride will be present in an amount of from about 0.1 to 3.0% by weight of the total weight of the oral composition, preferably from about 0.4 to 2.5% by weight. A typical toothpaste composition may contain 2% by weight while a typical mouthwash composition may contain 0.5% by weight of the guar hydroxypropyltrimonium chloride.

Brief Summary Text (25):

The concentration of cetylpyridinium chloride and guar hydroxypropyltrimonium chloride will depend, in part, on the form of the composition (i.e. a solution such as mouthwash or gargle or a semi-solid such as a toothpaste, lozenge, and chewing gum) which is used to deliver cetylpyridinium chloride to the gingiva/mucosal tissue and/or the tooth surfaces. For example, solutions containing cetylpyridinium chloride are generally more efficient in contacting the tissue and tooth surfaces than semi-solid compositions and therefore may require a lower concentration of the antibacterial agent.

Brief Summary Text (26):

Orally acceptable carriers are typically present in the oral composition of the present invention. The term "orally acceptable carrier" as used herein is meant to include one or more compatible solid or liquid filler diluents or encapsulating substances which are suitable for topical, oral administration. It is understood that the components of the composition are preferably capable of being commingled without interaction in a manner which would substantially reduce the composition's stability and/or efficacy for reducing the presence of microorganisms in the oral

cavity according to the compositions and methods of the present invention. The orally acceptable carriers of the present invention can include the usual and conventional components of toothpastes (including pastes, gels and gels for subgingival application), mouthwashes, mouth sprays, gargles, denture cleansers, chewing gums, and lozenges (including breath mints). Such orally acceptable carriers include, but not limited to, water, oils, organic solvents, saline, alcohols such as ethanol, glycerin, waxes, soft paraffin bases, petrolatum, lanolin, mixtures thereof and the like. The orally acceptable carrier will vary according to the form of the composition and is typically present in an amount of from about 20 to 99% by weight. A typical toothpaste composition may contain 15% glycerol and 20% water. A typical mouthwash composition may contain 94% of the sum of water, alcohol, and glycerin.

Brief Summary Text (28):

Where the oral composition of the present invention is a gel or paste such as in a toothpaste (dentifrice), dental cream dentifrices or gel dentifrices, an orally acceptable carrier, including a water-phase with humectant which is preferably glycerin or sorbitol or an alkylene glycol such as polyethylene glycol or propylene glycol may be present. Such gel or paste compositions typically further contain a natural or synthetic thickener or gelling agent.

Brief Summary Text (29):

Where the oral composition of the present invention is a solution or a liquid such as a mouthwash, the orally acceptable carrier is typically a water-alcohol mixture.

Brief Summary Text (31):

The mouthwash form of the oral composition may be suitably prepared by mixing the appropriate components thereof. Dentifrices are prepared in a similar manner with the addition, typically, of a thickener and a polishing agent.

Brief Summary Text (34):

The present oral composition may optionally contain a sweetening agent for masking the objectionable taste often associated with cetylpyridinium chloride and improving the organoleptic properties of the oral composition. Suitable sweetening agents include high potency sweeteners such as cyclamate, saccharine, acesulfame, and sucralose, and the orally acceptable salts thereof, and bulk sweeteners such as xylitol, sorbitol, erythritol. Additional sweeteners such as sucrose, lactose, maltose, glucose, and fructose may also be used, but are not desirable due to their cariogenic potential.

Brief Summary Text (36):

Antibacterial agents other than cetylpyridinium chloride may be optionally present in the oral compositions of the present invention. Such agents may include, but not limited to, chlorhexidine gluconate; benzalkonium chloride; benzethonium chloride; domiphen bromide; zinc salts such as zinc chloride, citrate or gluconate; stannous salts such as stannous chloride and fluoride; triclosan; sanguinarine chloride; and essential oils such as eucalyptol, thymol, menthol and eugenol. If present, the additional antibacterial agents generally comprise up to about 2% by weight, preferably from about 0.1 to 2% by weight of the composition of the present invention.

Brief Summary Text (39):

Abrasive agents include certain phosphates such as sodium metaphosphate, potassium metaphosphate, tricalcium phosphate, dicalcium phosphate dihydrate, anhydrous dicalcium phosphate, calcium pyrophosphate, zinc orthophosphate, alumina, hydrated alumina, aluminum silicate, bentonite, calcium carbonate, sodium bicarbonate and the like. The abrasive agents are typically present in an amount of from about 0.5 to 70% by weight, preferably from about 5 to 40% by weight based on the total weight of the composition.

Brief Summary Text (40):

Fluoride-supplying compounds include inorganic fluoride salts, such as soluble alkali metal, alkaline earth metal, and heavy metal fluoride salts, for example, sodium fluoride, potassium fluoride, ammonium fluoride, lead fluoride, copper fluoride, zinc fluoride, tin fluoride, barium fluoride, sodium fluorosilicate, ammonium fluorosilicate, sodium fluorozirconate, sodium monofluorophosphate, and the

like. The fluoride-supplying compound is typically present in an amount sufficient to deliver from about 50 to 5,000 ppm, preferably from about 100 to 2,500 ppm, more preferably from about 1,000 to 1,200 ppm of available fluoride based on the composition.

Brief Summary Text (44):

In accordance with a preferred aspect of the present invention, a cetylpyridinium chloride-containing toothpaste composition containing guar hydroxypropyltrimonium chloride alone or in combination with a sweetening agent, especially sodium saccharin dihydrate provides significant availability of the antibacterial agent to bind to tooth surfaces.

Brief Summary Text (45):

A safe and effective amount of the compositions of the present invention may be topically applied in several conventional ways to the mucosal tissue of the oral cavity, to the gingival tissue of the oral cavity, and/or to the tooth surface, for reducing the levels of undesirable oral microorganisms residing thereon. For example, the gingival or mucosal tissue may be rinsed with a solution (e.g., mouthwash, mouth spray) containing cetylpyridinium chloride and guar hydroxypropyltrimonium chloride, or if cetylpyridinium chloride and guar hydroxypropyltrimonium chloride are included in a dentifrice (e.g., toothpaste, tooth gel, or tooth powder), the gingival or mucosal tissue is bathed in the liquid and/or in the lather generated by brushing of the teeth.

Brief Summary Text (46):

Other non-limiting examples include applying a non-abrasive gel or paste, which contains cetylpyridinium chloride and guar hydroxypropyltrimonium chloride, directly to the gingival/mucosal tissue or to the tooth surface with or without an oral care implement; chewing gum that contains cetylpyridinium chloride and guar hydroxypropyltrimonium chloride; chewing or sucking on a breath tablet or lozenge which contains cetylpyridinium chloride and guar hydroxypropyltrimonium chloride. Preferred methods of using compositions of the present invention include applying cetylpyridinium chloride and guar hydroxypropyltrimonium chloride to the gingival/mucosal tissue and/or the tooth surface via rinsing with a mouthwash solution and via brushing with a dentifrice. Other methods of applying cetylpyridinium chloride and guar hydroxypropyltrimonium chloride to the gingival/mucosal tissue and tooth surfaces are apparent to those skilled in the art.

Detailed Description Text (2):

Effect of Guar Hydroxypropyltrimonium Chloride on Adsorption of Cetylpyridinium Chloride in the Presence of a Sweetening Agent

Detailed Description Text (3):

In vitro studies of cetylpyridinium chloride absorption were performed with the initial step of formulating a model oral surface. Disks of hydroxyapatite (HAP, calcium phosphate hydroxide, $\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_2$) measuring 0.5 inch diameter and 0.04 inch thick were obtained from Clarkson Chromatography Products (Williamsport, Pa.). The disks were hydrated in deionized water for one hour and then allowed to air dry.

Detailed Description Text (5):

Cetylpyridinium chloride adsorption was tested by soaking the disks in a test solution. The compositions of the test solutions are shown and listed in Table 1. Guar hydroxypropyltrimonium chloride was obtained from Hercules Incorporated, Wilmington, Del., which is marketed under the trade name "N-Hance 3215". The disks were each soaked in 5 mL of the test solution for 10 minutes in a polystyrene petri dish (35 mm diameter.times.10 mm deep, from Becton Dickinson). The disks were then removed, rinsed for 3 seconds with deionized water on each side with a wash bottle.

Detailed Description Text (6):

Adsorbed cetylpyridinium chloride was extracted by soaking the disks in a solution used as a mobile phase for cetylpyridinium chloride detection in liquid chromatography. The extractant solution was composed of 60 parts of a 20 mM phosphate buffer and 40 parts methanol, in which was dissolved 30 mM

cetyltrimethylammonium bromide (CTAB). The disks were soaked in 5 mL of the extractant solution for 2 hours. The extractant was then analyzed for cetylpyridinium chloride using high-pressure liquid chromatography.

Detailed Description Text (7):

The composition of each test solution and the results of cetylpyridinium chloride absorption tests are shown in Table 1.

Detailed Description Text (8):

As shown in Table 1, the addition of guar hydroxypropyltrimonium chloride to a solution containing saccharin increased the adsorption of cetylpyridinium chloride by about a factor of at least two as compared to the solution containing only cetylpyridinium chloride and saccharin.

Detailed Description Text (10):

Effect of Guar Hydroxypropyltrimonium Chloride on Adsorption of Cetylpyridinium Chloride in the Presence of an Emollient

Detailed Description Text (11):

In vitro studies of cetylpyridinium chloride adsorption were performed with the initial step of formulating a model oral surface. Disks of hydroxyapatite (HAP, calcium phosphate hydroxide, Ca.sub.10 (PO.sub.4).sub.6 (OH).sub.2) measuring 0.5 inch diameter and 0.04 inch thick were obtained from Clarkson Chromatography Products (Williamsport, Pa.). The disks were hydrated in deionized water for one hour and then allowed to air dry.

Detailed Description Text (13):

Cetylpyridinium chloride absorption was tested by soaking the disks in a test solution. The test solutions are shown and listed in Table 2. Guar hydroxypropyltrimonium chloride was obtained from Hercules Incorporated, Wilmington, Del., which is marketed under the trade name "N-Hance 3215". Eldew CL-301 or cholesteryl/behanyl/octyldecyl lauroyl glutamate, is an emollient derived from L-glutamic acid, lauric acid and three alcohols (cholesterol, 2-octyldodecanol, and behenol), and is marketed and manufactured by Ajinomoto Inc. (Tokyo, Japan). The disks were each soaked in 5 mL of the test solution for 10 minutes in a polystyrene petri dish (35 mm diameter.times.10 mm deep, from Becton Dickson). The disks were then removed, rinsed for 3 seconds with deionized water on each side with a wash bottle.

Detailed Description Text (14):

Adsorbed cetylpyridinium chloride was extracted by soaking the disks in a solution used as a mobile phase for cetylpyridinium chloride detection in liquid chromatography. The extractant solution was composed of 60 parts of a 20 mM phosphate buffer and 40 parts methanol, in which was dissolved 30 mM cetyltrimethylammonium bromide (CTAB). The disks were soaked in 5 mL of the extractant solution for 2 hours. The extractant was then analyzed for cetylpyridinium chloride using high-pressure liquid chromatography.

Detailed Description Text (15):

The composition of each test solution and the results of cetylpyridinium chloride adsorption tests are shown in Table 2.

Detailed Description Text (16):

As shown in Table 2, the addition of guar hydroxypropyltrimonium chloride to a solution containing Eldew CL-301 increased the adsorption of cetylpyridinium chloride by about a factor of at least four as compared to the solution containing only cetylpyridinium chloride and Eldew CL-301.

CLAIMS:

1. An oral composition comprising:

- a) an antibacterial effective amount of cetylpyridinium chloride;
- b) an effective amount of guar hydroxypropyltrimonium chloride, sufficient to bind

to compounds which undesirably bind to cetylpyridinium chloride thereby enabling the cetylpyridinium chloride to effectively bind to tooth surfaces and perform an antibacterial function; and

c) an orally acceptable carrier.

3. The oral composition of claim 1, wherein the oral composition is in a form selected from the group consisting of a mouthwash, a dentifrice, a chewing gum, and a lozenge.

4. The oral composition of claim 1, the amount of cetylpyridinium chloride is present from about 0.01 to 1.0% by weight based on the total weight of the oral composition.

6. The oral composition of claim 3 wherein the dentifrice is a toothpaste.

8. The oral composition of claim 7 wherein the abrasive agents are selected from the group consisting of sodium metaphosphate, potassium metaphosphate, tricalcium phosphate, dicalcium phosphate dihydrate, anhydrous dicalcium phosphate, calcium pyrophosphate, zinc orthophosphate, alumina, hydrated alumina, aluminum silicate, bentonite, calcium carbonate, and sodium bicarbonate.